



**(12)**

**EUROPEAN PATENT APPLICATION**

**(21) Application number : 93306886.8**

**(51) Int. Cl.<sup>5</sup> : C07H 19/048, C07H 19/06**

**(22) Date of filing : 31.08.93**

**(30) Priority : 01.09.92 US 938791**

**(43) Date of publication of application :  
16.03.94 Bulletin 94/11**

**(84) Designated Contracting States :  
AT BE CH DE DK ES FR GB GR IE IT LI LU NL  
PT SE**

**(71) Applicant : ELI LILLY AND COMPANY  
Lilly Corporate Center  
Indianapolis Indiana 46285 (US)**

**(72) Inventor : Britton, Thomas Charles  
4700 Royal Oak Lane  
Carmel, Indiana 46033 (US)  
Inventor : Le Toumeau, Michael Edward  
7566 Augusta Court  
Indianapolis, Indiana 46268 (US)**

**(74) Representative : Tapping, Kenneth George et  
al  
Lilly Industries Limited Patent Department Erl  
Wood Manor  
Windlesham Surrey, GU20 6PH (GB)**

**(54) A process for anomerizing nucleosides.**

**(57) A process for increasing the amount of beta-anomer nucleoside from an alpha-anomer nucleoside or undesired anomeric mixture of nucleosides by contacting the anomer or anomeric mixture with a hydroxide base in an organic solvent.**

The invention relates to the field of pharmaceutical and organic chemistry and pertains to a process for anomerizing nucleosides.

Processes for preparing nucleosides frequently result in a mixture of alpha and beta nucleoside anomers. These nucleoside anomers are typically separated by a physical means such as crystallization or chromatography. Most often, the desired biological activity of a nucleoside resides predominantly in a single anomer of an anomeric mixture. However, the amount of a specified nucleoside anomer recoverable from an anomeric mixture by the above mentioned separation methods is often substantially less than that originally present in the anomeric mixture. Such low recoveries are generally due to interference from increased proportions of the unwanted anomer as the separation proceeds. Beta nucleoside anomers are useful and important as pharmacologically active compounds. Anomerization provides a way of increasing the amount of a desired nucleoside anomer over that originally present in an anomeric mixture. When used in conjunction with the aforementioned separation methods, anomerization can afford substantially improved overall recoveries of a desired nucleoside anomer.

Nucleoside anomerization has been accomplished by photoirradiation in water, see R. A. Sanchez, *et al.*, *J. Mol. Biol.*, **47**, 531-543 (1970); and with bromine, see H. Quelo, *et al.*, *C. R. Acad. Sci., Ser. C*, **275**, 1137-1140 (1972).

J. cadet, *et al.*, describe nucleoside anomerization in *Nucleic Acid Hydrolysis I. Isomerization and Anomerization of Pyrimidic Deoxyribonucleosides in an Acidic Medium.*, *J. Amer. Chem. Soc.*, **96**:20, 6517-6519 (1974) which involves contacting thymidine and 2'-deoxyuridine nucleosides with 2 M HClO<sub>4</sub> at 90°C to make α- and β-furanosidic and pyranosidic anomers.

Yamaguchi, T., *et al.*, in *Synthetic Nucleosides and Nucleotides. XXI. On the Synthesis and Biological Evaluations of 2'-Deoxy-α-D-ribofuranosyl Nucleosides and Nucleotides*, *Chem. Pharm. Bull.*, **32**(4), 1441-1450 (1984) describe anomerizing β-3', 5'-di-O-p-toluoyl-2'-deoxythymidine and β-N<sup>4</sup>-benzoyl-2'-deoxycytidine with bis(trimethylsilyl) acetamide and trimethylsilyltrifluoromethanesulfonate in dry acetonitrile at 70°C.

Nucleoside anomerization employing protic acids or Lewis acids have been applied to a wide variety of nucleosides and include for example: 2 M HCl, see F. Seela and H. D. Winkler, *Carbohydrate Research*, **118**, 29-53 (1983); 1 M HBr, see J. Cadet, *Tetrahedron Lett.*, 867-870 (1974); and Na/HOAc, see J. Matulic-Adamic, *et al.*, *J. Chem. Soc.*, 2681-2686 (1988).

Base catalyzed anomerization has also been reported. For example, Armstrong, V.W., *et al.*, in *The Base Catalyzed Anomerization of β-5-Formyluridine; Crystal and Molecular Structure of α-5-Formyluridine*, *Nucleic Acid Res.*, **3**, 1791 (1976) describe the treatment of β-5-formyluridine with 1:1 4 N aqueous NaOH:MeOH at room temperature which affords an anomerically mixed product. However, uridine and 5-bromouridine are not anomerized by this process since they lack the 5-formyl group on the nucleoside substrate. I. Hideo, *et al.*, *Synthesis of 5-Alkyl and 5-Acyl-uridines via 6-Mercaptouridine (Nucleosides and Nucleotides XVII)*, *Heterocycles*, **8**, 427-432 (1977) describe the anomerization of 2',3'-O-isopropylidene-5-acetyl-α-uridine with 2 N sodium hydroxide. As can be seen, base catalyzed anomerization has been limited to pyrimidine nucleosides having electron-withdrawing substituents (e.g. formyl or acetyl groups) at the C-5 position of the heterocyclic portion of the nucleoside.

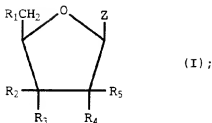
An object of the present invention is to provide a base catalyzed process for anomerizing nucleosides.

Another object of this invention is to provide a base catalyzed process for anomerizing 2'-deoxy-2',2'-difluoro-nucleosides.

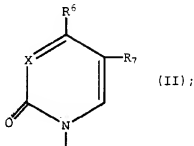
Another object of this invention is to provide a base catalyzed process for anomerizing alpha-anomer-enriched nucleosides free of the disadvantages and limitations found in the prior art.

Another object of this invention is to provide a base catalyzed process for anomerizing beta-anomer-enriched nucleosides free of the disadvantages and limitations found in the prior art.

According to the present invention there is provided a process for increasing the amount of beta-anomer nucleoside of the formula



wherein  $R_1$  is selected from the group consisting of hydrogen, lower alkyl, fluoro, azide, hydroxy, and OB where B is a lower alkyl, or base-stable hydroxy protecting group;  $R_2$  is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided  $R_3$  cannot be fluoro, azide, or hydroxy), and OB where B is as defined above;  $R_3$  is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided  $R_2$  cannot be fluoro, azide or hydroxy), and OB where B is as defined above;  $R_4$  is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided  $R_5$  cannot be fluoro, azide or hydroxy), and OB where B is as defined above;  $R_5$  is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided  $R_4$  cannot be fluoro, azide or hydroxy), and OB where B is as defined above; and Z is a nucleobase of the formula



wherein X is selected from N and CR<sub>6</sub> where  $R_6$  is hydrogen or lower alkyl;  $R_6$  is selected from the group consisting of amino, lower alkyl amino, di(lower alkyl) amino, acyl amino, and N-acyl lower alkyl amino; and  $R_7$  is selected from the group consisting of hydrogen, lower alkyl, fluoro and lower alkenyl; in an alpha-anomer enriched nucleoside over that originally present; comprising contacting an alpha-anomer enriched nucleoside of formula (I) with a hydroxide base and an organic solvent.

Throughout this document, all temperatures are in degrees Celsius and all proportions, percentages and the like, are in weight units. Mixtures of solvents are in volume units, except where otherwise indicated. Anomeric mixtures are expressed as a weight/weight ratio. The phrase "anomer enriched" alone or in combination refers to an anomeric mixture wherein the anomeric ratio differs from the equilibrium anomeric ratio and includes substantially pure anomers. The term "lower alkyl" alone or in combination refers to straight, cyclic and branched chain aliphatic hydrocarbon groups which preferably contain up to 7 carbon atoms such as methyl, ethyl, n-propyl, isopropyl, n-butyl, t-butyl, n-pentyl, n-hexyl, 3-methylpentyl groups and the like. The term "aryl" alone or in combination refers to carbocyclic or heterocyclic groups such as phenyl, naphthyl, thienyl and substituted derivatives thereof. The term "acyl" alone or in combination refers to the general formula ACO; wherein A is lower alkyl or aryl. The term "lower alkenyl" refers to an unsaturated hydrocarbon group containing up to 7 carbon atoms and having one or two carbon double bonds. The phrase "base-stable hydroxy protecting group" refers to hydroxy protecting groups stable under basic conditions as described in Chapter 3 of Protective Groups in Organic Chemistry, McOmie Ed., Plenum Press, New York (1973) and Chapter 2 of Protective Groups in Organic Synthesis, Green, John, J. Wiley and Sons, New York (1981) such as benzyloxymethyl, methoxymethyl, 2-tetrahydropyranyl, benzyl, p-methoxybenzyl and trityl; and where the nucleoside contains a cis-2',3'-diol derivative the base-stable hydroxy protecting group includes acetone, benzylidene and p-methoxybenzylidene.

The present process is carried out by contacting an alpha-anomer enriched nucleoside of formula (I) with a hydroxide base in an organic solvent. The process promotes the stereoconversion of nucleosides by inverting the absolute configuration at the C-1' position of the nucleoside. While not wishing to be bound by theory, it is believed that this inversion is achieved by the action of the hydroxide base, hydroxide base concentration,

solvent, and reaction temperature employed.

The present process increases the amount of beta-anomer nucleoside present in an anomeric mixture of unprotected nucleosides and nucleosides typically unreactive to acid catalyzed anomerization processes.

A preferred embodiment of the present process, employs alpha-anomer enriched nucleosides having an anomeric ratio ranging from at least 10:90 alpha to beta up to substantially pure (about 100:0) alpha anomer; and more preferably ranging from about 50:50 alpha to beta up to substantially pure alpha-anomer.

A particularly preferred embodiment of the present process, increases the amount of 2',2'-difluoro-2'-deoxy-beta-anomer nucleoside in anomeric mixture and employs an alpha-anomer enriched 2',2'-difluoro-2'-deoxy nucleoside having an anomeric ratio ranging from at least 75:25 alpha to beta to substantially pure alpha-anomer.

Hydroxide bases useful in the present process include alkali metal hydroxides such as lithium hydroxide, sodium hydroxide, potassium hydroxide and cesium hydroxide monohydrate; quaternary ammonium hydroxide bases such as benzyltrimethylammonium hydroxide and tetramethylammonium hydroxide; and alkaline earth metal hydroxides; most preferred are alkali metal hydroxides such as potassium hydroxide and cesium hydroxide monohydrate. The amount of hydroxide base employed in the present process ranges from about 2 molar equivalents to about 40 molar equivalents; however, from about 2.5 molar equivalents to about 5 molar equivalents is preferred.

It has been found that the rate of anomerization exhibits a third order dependency on the hydroxide base concentration. Therefore, the of hydroxide base concentration preferably ranges from about 0.5 molar to about 5 molar and more preferably ranges from about 2 molar to about 4 molar.

Solvents useful in the present process are C<sub>1</sub>-C<sub>7</sub> alcohols such as methanol, ethanol, 2-methoxyethanol, and mixtures thereof; preferred is methanol.

The reaction time is a function of the reactivity of the nucleoside, the hydroxide base, the hydroxide base concentration and the reaction temperature employed. The present process is preferably carried out at temperatures ranging from room temperatures to about 120°C; more preferably from about 40°C to about 120°C; and most preferably from about 40°C to about 80°C. The present process is carried out in about 1/2 hour to 5 days.

The present process shifts the anomeric ratio of an anomer enriched nucleoside towards its equilibrium anomeric ratio. The equilibrium anomeric ratio differs for each nucleoside, however, for anomeric mixtures of 1-(2'-deoxy-2',2'-difluororibofuranosyl)-4-aminopyrimidin-2-one the equilibrium anomeric ratio is approximately 60 (β):40(α). It should be noted that the reaction rate decreases substantially as the equilibrium anomeric ratio is approached. Therefore, the present process is carried out, in either a batch, semi-batch or continuous mode, so that it may be stopped prior to reaching the equilibrium anomeric ratio in order to avoid yield losses due to competing reactions, e.g. hydrolysis.

When the present process is carried out in the presence of water, the potential for producing hydrolysis products, e.g. the conversion of the cytosine nucleobase to a uracil nucleobase, increases. However, when the present process is carried out in substantially anhydrous organic solvents, the hydrolysis reaction is suppressed and a substantially higher yield of the anomerization product results. Therefore, the amount of water employed in the present process is substantially zero.

The process may be monitored by withdrawing aliquots at various times over the course of the reaction, quenching the aliquots with acid, diluting the aliquots with an appropriate volume with water, and assaying the aliquots by high pressure liquid chromatography (HPLC) to determine the anomer ratio of the nucleosides present.

Once the desired anomer ratio has been achieved, the resulting solution is acidified, for example, by adding an acid such as hydrochloric acid, or neutralized, depending on the nucleoside employed.

The desired nucleoside anomer may be isolated by standard separation techniques such as crystallization or chromatography.

#### Example 1

Anomerization of 1-(2'-deoxy-2',2'-difluoro-α-D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro-β-D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Anhydrous Lithium Hydroxide in Methanol.

A solution of 1.50 g (5.70 mmol) of 1 in 6.0 ml of anhydrous methanol was treated with 410 mg (17.1 mmol; 3.0 eq.) of anhydrous lithium hydroxide and the resulting mixture was heated to reflux under dry nitrogen. Reaction aliquots (0.100 ml, 1.40 % of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs.)	1	2	1:2
1	0.33	99.6	0.4	100.0
2	24.50	64	18	79:21
3	51.25	42	24	64:36
4	71.50	34	24	58:42
5	94.75	27	24	53:47

**Method A:** Column: 25 cm x 4.6 mm Zorbax RX reverse phase. Flow rate: 1.2 ml/min. Solvent A: methanol. Solvent B: 0.1 M pH 3 phosphate buffer. Gradient program: 0 - 8.0 minutes isocratic 3/97 of A/B; 8.0 - 13.0 minutes linear gradient from 3/97 of A/B to 50/50 of A/B; 13.0 - 18.0 minutes isocratic 50/50 of A/B; 18.0 - 23.0 minutes linear gradient from 50/50 of A/B to 3/97 of A/B. The peak areas of 1 ( $t_r$  = 4.9 minutes) and 2 ( $t_r$  = 7.2 minutes) were compared to an external standard containing known quantities of authentic samples to provide the yields of each.

### Example 2

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Anhydrous Sodium Hydroxide in Methanol

A solution of anhydrous sodium hydroxide in methanol was prepared by adding 6.0 ml of anhydrous methanol, stirred at 25°C under dry nitrogen, to 393 mg (17.1 mmol, 3.0 eq.) of sodium metal. When the metal dissolved, water (306  $\mu$ l, 17.0 mmol, 3.0 eq.) was added. To the above solution was added 1.50 g (5.70 mmol) of 1, and the resulting mixture was heated to reflux. Reaction aliquots (0.100 ml, 1.36% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs.)	1	2	1:2
1	.33	99	0.7	>99:1
2	1.50	92	6	94:6
3	18.50	40	30	57:43
4	23.00	34	30	53:47
5	25.75	31	30	51:49
6	90.75	13	18	43:57

### Example 3

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Potassium Hydroxide in Ethanol

A solution of 1.50 g (5.70 mmol) of 1 in 6.0 ml of absolute ethanol was treated with 1.10 g (17.1 mmol; 3.0 eq.) of 86 percent potassium hydroxide and the resulting mixture was heated to 76°C-77°C under dry nitrogen. Reaction aliquots (0.100 ml, 1.26% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs. )	1	2	1:2
1	.33	93	5	95:5
2	2.00	49	22	69:31
3	4.50	24	23	51:49
4	6.50	17	21	45:55
5	24.33	4	6	39:61
6	29.00	3	5	39:61

**Example 4**

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Barium Hydroxide in Methanol

A mixture of 0.60 g (2.28 mmol; 1.0 eq.) of 1, 0.62 g (3.42 mmol; 1.5 eq.) of 95 percent barium hydroxide and 4.4 ml of anhydrous methanol was stirred and heated at reflux for 28 hours. The resulting mixture was cooled to 0°C, quenched with 5.6 ml of 1 N HCl, and diluted to 250 ml with water. A 5.00 ml aliquot of the resulting tan solution was diluted to 100.0 ml with water, and assayed by HPLC (Method A). The yield of 1 and 2 and their anomeric ratio (1:2) is tabulated below:

% Yield	% Yield	Ratio
1	2	1:2
82	8	92:8

**Example 5**

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Cesium Hydroxide Monohydrate in Methanol

A mixture of 1.23 g (4.68 mmol) of 1, 2.36 g (14.05 mmol; 3.0 eq.) of cesium hydroxide monohydrate, and 4.93 ml of anhydrous methanol was heated to reflux under dry nitrogen. Reaction aliquots (0.100 ml, 1.59% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

Aliquot	Elapsed Time (Hrs.)	% Yield 1	% Yield 2	Ratio 1:2
1	.33	97	3	97:3
2	2.50	73	21	78:22
3	4.50	58	31	65:35
4	7.00	48	37	56:44
5	24.00	23	32	41:59

## Example 6

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Potassium Hydroxide in 2-Methoxyethanol

A mixture of 1.50 g (5.70 mmol) of 1, 1.10 g (16.9 mmol; 3.0 eq.) of 86 percent potassium hydroxide, and 6.0 ml of 2-methoxyethanol was heated to 76°C under dry nitrogen. Reaction aliquots (0.100 ml, 1.26% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs. )	1	2	1:2
1	.42	93	4	96:4
2	2.08	62	17	78:22
3	4.58	38	23	62:38
4	6.58	29	24	54:46
5	24.50	6	12	38:64
6	29.08	5	10	35:65

## Example 7

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Potassium Hydroxide in Methanol

A mixture of 750 mg (2.85 mmol) of 1, 558 mg (8.55 mmol; 3.0 eq.) of 86 percent potassium hydroxide, and 3.4 ml of anhydrous methanol was heated to reflux under dry nitrogen. Reaction aliquots (0.100 ml, 2.58% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs.)	1	2	1:2
1	.33	99	1	99:1
2	2.17	88	12	88:12
3	3.50	78	18	81:19
4	4.92	70	22	76:24
5	24.00	29	34	46:54
6	29.00	27	34	44:56
7	47.25	21	29	42:58

## Example 8

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Potassium Hydroxide in Methanol

A mixture of 1.50 g (5.70 mmol) of 1, 1.10 g (16.9 mmol; 3.0 eq.) of 86 percent potassium hydroxide and 4.4 ml of anhydrous methanol was heated to 55°C under dry nitrogen. Reaction aliquots (0.100 ml, 1.72% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

below:

	Elapsed	% Yield	% Yield	Ratio
<u>Aliquot</u>	<u>Time (Hrs.)</u>	<u>1</u>	<u>2</u>	<u>1:2</u>
1	.33	99	1	99:1
2	4.17	87	11	99:11
3	24.50	50	15	61:40
4	27.58	49	35	58:42
5	45.25	37	38	50:50

#### Example 9

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Benzyltrimethylammonium Hydroxide in Methanol

Three identical mixtures of 250 mg (0.95 mmol) of 1 and 1.3 ml (2.85 mmol, 3.0 eq.) of N-benzyltrimethylammonium hydroxide (40% by weight in methanol) were heated at reflux under dry nitrogen for the times indicated below. The resulting solutions (1-3) were cooled to 25°C and each was quenched by the adding 10 ml of 1.0 N HCl, diluted to 1 L with water, and assayed by HPLC (Method A). The yields of 1 and 2 and their anomer ratios (1:2) are tabulated below:

	Reflux	% Yield	% Yield	Ratio
Solution	Time (Hrs. )	1	2	1:2
1	3.0	65	15	81:19
2	5.5	52	22	71:29
3	8.0	35	24	59:41

#### Comparative Example 10

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one Hydrochloride (1 · HCl) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Aqueous Sodium Hydroxide

This example illustrates the effect water has on the nucleoside anomer yield. A solution of 160 mg (0.53 mmol; 1.0 eq.) of 1 · HCl in 40 ml of 2.0 N aqueous NaOH (80 mmol, 150 eq.) was heated at 60°C. Reaction aliquots (4.00 ml, 10.0% of the total) were withdrawn at the indicated times, quenched with 10 ml of 1 N HCl, diluted to 50.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomer ratios (1:2) are tabulated below:



	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs. )	1	2	1:2
1	1.0	68	10	87:13
2	3.0	35	13	73:27
3	6.0	12	8	62:38
4	24.0	0	0	---

**Example 11**

Anomerization of a crude 81:19 mixture of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (1) and 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) with Potassium Hydroxide in Methanol

The selective crystallization of 2 from a crude aqueous mixture of 1 and 2 having an anomeric ratio (1:2) of 85:35 provided a mother liquor having an anomeric ratio of 81:19. On evaporating the liquor in vacuo 38.14 g of residue was obtained which was found by HPLC analysis to contain 18.32 g (0.070 moles) of total nucleoside (1 and 2). A solution of the above residue, 13.7 g (0.210 moles; 3.0 eq.) of 86 percent potassium hydroxide and 120 ml of methanol was heated at reflux under dry nitrogen. After 8.25 hours, an additional 2.3 g (0.035 moles) of 86 percent potassium hydroxide was added over a 10 minute period. Reaction aliquots (0.100 ml, 0.0645% of the total) were withdrawn at the times indicated below, quenched with 5 ml 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed	% Yield	% Yield	Ratio
Aliquot	Time (Hrs.)	1	2	1:2
1	.42	81	19	81:19
2	7.50	70	24	75:25
3	8.25	70	25	74:26
4	27.50	49	36	58:42

**Example 12**

Anomerization of 1-(2'-deoxy-2',2'-difluoro- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one Hydrochloride (1 ·HCl) to 1-(2'-deoxy-2',2'-difluoro- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (2) Potassium Hydroxide in Methanol

A mixture of 1.60 g (5.34 mmol) of 1·HCl, 1.40 g (21.5 mmol; 4.0 eq.) of 86 percent potassium hydroxide, and 7.5 ml of anhydrous methanol was heated at reflux under dry nitrogen. Reaction aliquots (0.135 ml, 1.47% of the total) were withdrawn at the times indicated below, quenched with 5 ml of 1 N HCl, diluted to 100.0 ml with water and assayed by HPLC (Method A). The yields of 1 and 2 and their anomeric ratios (1:2) are tabulated below:

	Elapsed %	% Yield	% Yield	Ratio
Aliquot	Time (Hrs)	1	2	1:2
1	1.08	95	4	96:4
2	4.33	82	14	85:15
3	7.08	73	21	77:23
4	23.08	45	38	55:45
5	29.58	39	40	50:50

After refluxing for 30 hours, the reaction mixture was cooled in an ice bath and acidified by the dropwise addition of 1.5 ml of concentrated HCl. The resulting mixture was filtered to remove the precipitated salts and the filter cake was washed with methanol (3 x 5 ml portions). The filtrate was then evaporated in vacuo and the residue dissolved in 7 ml of water. The pH of the aqueous solution was adjusted to 7 with aqueous potassium hydroxide and the solution was concentrated in vacuo until crystallization ensued. Upon cooling for 16 hours at 5°C-10°C 328 mg (after air drying) of off-white precipitate were obtained and shown by <sup>1</sup>H NMR and HPLC analysis (Method A) to be 83.9 percent of 2 and contained 1 percent total non-volatile impurities, for a 21 percent isolated yield of 2.

#### Example 13

Anomerization of 1-(2'-deoxy- $\alpha$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (5) to 1-(2'-deoxy- $\beta$ -D-ribofuranosyl)-4-aminopyrimidin-2-one (6) with Potassium Hydroxide in Methanol

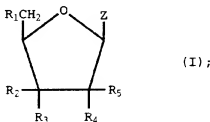
A mixture of 1.14 g (5.0 mmol) of 5, 7.5 ml of methanol and 990 mg (15.0 mmoles; 3.0 eq.) of 85 percent potassium hydroxide was heated at reflux. Reaction aliquots (80  $\mu$ L, 0.929% of the total) were withdrawn at the times indicated below, quenched with 25 ml of 0.05 M pH 3 phosphate buffer and diluted to 100.0 ml with water and assayed by HPLC (Method C). The yields of 5 and 6 and their anomeric ratios (5:6) are tabulated below:

	Elapsed Time			
Aliquot	(Hrs. )	% Yield	% Yield	Ratio
1	0.5	99.5	0.5	99.5:0.5
2	5.0	92.8	4.6	95:5
3	22.0	76.1	15.2	83:17
4	28.0	72.0	17.8	80:20
5	46.0	61.2	22.5	73:27
6	52.5	59.1	23.8	71:29
7	71.5	50.7	24.7	67:33
8	101.5	43.2	24.7	64:36
9	124.0	40.9	24.9	62:38

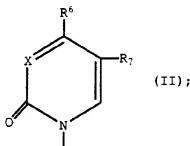
Method C: 25 cm X 4.6 mm Apex ODS 5  $\mu$  column. Flow rate: 0.8 ml/minute. Solvent A: methanol. Solvent B: 0.05 M pH 3 phosphate buffer. Gradient: 0- 10 minutes isocratic 100% of B; 10-15 minutes linear gradient from 100% of B to 50/50 of A/B; 15-19 minutes, isocratic 50/50 of A/B; 19-23 minutes linear gradient from 50/50 of A/B to 100% of B. The peak areas of 5 ( $t_r$  = 6.6 minutes) and 6 ( $t_r$  = 8.1 minutes) were compared to an external standard containing known quantities of authentic samples to provide the yields of each.

## Claims

1. A process for increasing the amount of beta- anomer nucleoside of the formula



15 wherein R<sub>1</sub> is selected from the group consisting of hydrogen, lower alkyl, fluoro, azide, hydroxy, and OB where B is a lower alkyl or base-stable hydroxy protecting group; R<sub>2</sub> is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided R<sub>3</sub> cannot be fluoro, azide, or hydroxy), and OB where B is as defined above; R<sub>3</sub> is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided R<sub>2</sub> cannot be fluoro, azide or hydroxy), and OB where B is as defined above; R<sub>4</sub> is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided R<sub>5</sub> cannot be fluoro, azide or hydroxy), and OB where B is as defined above; R<sub>5</sub> is selected from the group consisting of hydrogen, azide, lower alkyl, fluoro, hydroxy (provided R<sub>4</sub> cannot be fluoro, azide or hydroxy), and OB where B is as defined above; and Z is a nucleobase of the formula



wherein X is selected from N and CR<sub>8</sub> where R<sub>8</sub> is hydrogen or lower alkyl; R<sub>6</sub> is selected from the group consisting of amino, lower alkyl amino, di(lower alkyl) amino, acyl amino, and N-acyl lower alkyl amino; and R<sub>7</sub> is selected from the group consisting of hydrogen, lower alkyl, fluoro and lower alkenyl; in an alpha-anomer enriched nucleoside over that originally present; comprising contacting an alpha-anomer enriched nucleoside of formula (I) with a hydroxide base and an organic solvent.

2. The process of Claim 1 wherein the hydroxide base is selected from the group consisting of alkali metal hydroxides, alkaline earth metal hydroxides or quaternary ammonium hydroxides.
3. The process of Claim 2 wherein the hydroxide base is an alkali metal hydroxide selected from the group consisting of lithium hydroxide, potassium hydroxide, cesium hydroxide monohydrate or sodium hydroxide or an alkaline earth metal hydroxide such as barium hydroxide.
4. The process of Claim 3 wherein the alkali metal hydroxide is selected from potassium hydroxide, sodium hydroxide or cesium hydroxide monohydrate.
5. The process of Claim 1 wherein the amount of hydroxide base is from about 2 molar equivalents to about 40 molar equivalents.
6. The process of Claim 1 wherein the hydroxide base concentration is from about 0.5 molar to about 5 molar.
7. The process of Claim 1 wherein the solvent is selected from methanol, ethanol, 2-methoxyethanol, and mixtures thereof.

8. The process of Claim 7 wherein the solvent is a substantially anhydrous solvent.
9. The process of Claim 1 wherein the alpha anomer ratio is from about 10:90 alpha to beta to about 100:0 alpha to beta.
- 5 10. The process of Claim 1 wherein the process temperature is from room temperature to about 120°C.

10

15

20

25

30

35

40

45

50

55



European Patent  
Office

## EUROPEAN SEARCH REPORT

Application Number  
EP 93 30 6886

DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim
D, A	NUCLEIC ACIDS RESEARCH vol. 3, no. 7, 1976 pages 1791 - 1810 V.W.ARMSTRONG ET AL. 'The Base-catalysed Anomerisation of B-5-formyluridine; Crystal and Molecular Structure of alpha-5-formyluridine.' * abstract *	1
D, A	----- HETEROCYCLES vol. 8, 1977 pages 427 - 432 H. INOUE ET AL. 'Synthesis of 5-Alkyl- and 5-Acyl-Uridines via 6-Mercaptouridine (Nucleosides and Nucleotides, XVII).' * page 430, line 6 - line 12 * -----	1
		CLASSIFICATION OF THE APPLICATION (Int.Cl.5)
		C07H19/048 C07H19/06 A61K31/70
		TECHNICAL FIELDS SEARCHED (Int.Cl.5)
		C07H A61K
The present search report has been drawn up for all claims		
Place of search	Date of completion of the search	Examiner
THE HAGUE	17 December 1993	Scott, J
CATEGORY OF CITED DOCUMENTS		
<p>X : particularly relevant if taken alone  Y : particularly relevant if combined with another document of the same category  A : technological background  Q : non-written disclosure  P : intermediate document</p> <p>T : theory or principle underlying the invention  E : earlier patent document, but published on, or after the filing date  D : document cited in the application  L : document cited for other reasons</p> <p>A : number of the same patent family, corresponding document</p>		

EP 0 587 364 A1 (1993)